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     FILE 'HCAPLUS' ENTERED AT 16:11:32 ON 06 AUG 2009
L1       458707 S RNA OR RIBONUCLEIC
L2       705984 S ISOLATION OR PURIFICAITON OR EXTRACTION OR SEPARATION
L3       127166 S SOLID(W)(PHASE OR SUPPORT)
L4       364544 S LITHIUM
L5       173187 S ALKALI METAL
L6          233 S L1 AND L2 AND L3
L7           11 S L1 AND L2 AND L3 AND L4
L8            4 S L1 AND L2 AND L3 AND L5

     FILE 'STNGUIDE' ENTERED AT 16:11:41 ON 06 AUG 2009

     FILE 'HCAPLUS' ENTERED AT 16:12:31 ON 06 AUG 2009
L9      2168835 S ISOLATION OR PURIFICATION OR EXTRACTION OR SEPARATION OR ISOL
L10         473 S L1 AND L9 AND L3
L11          17 S L1 AND L9 AND L3 AND L4
L12           5 S L1 AND L9 AND L3 AND L5

     FILE 'STNGUIDE' ENTERED AT 16:12:35 ON 06 AUG 2009

     FILE 'HCAPLUS' ENTERED AT 16:12:49 ON 06 AUG 2009
L13          19 S L11 OR L12
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=> file hcaplus
COST IN U.S. DOLLARS                                SINCE FILE      TOTAL
                                                       ENTRY    SESSION
FULL ESTIMATED COST                                     0.88       0.88

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FILE COVERS 1907 - 6 Aug 2009  VOL 151 ISS 6
FILE LAST UPDATED: 4 Aug 2009  (20090804/ED)
REVISED CLASS FIELDS (/NCL) LAST RELOADED:  Jun 2009
USPTO MANUAL OF CLASSIFICATIONS THESAURUS ISSUE DATE:  Jun 2009

HCAplus now includes complete International Patent Classification (IPC)
reclassification data for the second quarter of 2009.

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The ALL, BIB, MAX, and STD display formats in the CA/CAplus family
of databases have been updated to include new citing references
information.  This enhancement may impact record import into
database management software.  For additional information, refer
to NEWS 22.

=> s RNA or ribonucleic

        377475 RNA
        195414 RIBONUCLEIC
L1      458707 RNA OR RIBONUCLEIC

=> s isolation or purificaiton or extraction or separation

        291282 ISOLATION
             6 PURIFICAITON
        193140 EXTRACTION
        242472 SEPARATION
L2      705984 ISOLATION OR PURIFICAITON OR EXTRACTION OR SEPARATION

=> s solid(w)(phase or support)

       1212483 SOLID
       2007475 PHASE
        573295 SUPPORT
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L3      127166 SOLID(W)(PHASE OR SUPPORT)

=> s lithium

L4      364544 LITHIUM

=> s alkali metal

        443198 ALKALI
       1946780 METAL
L5      173187 ALKALI METAL
                 (ALKALI(W)METAL)

=> s l1 and l2 and l3

L6         233 L1 AND L2 AND L3

=> s l1 and l2 and l3 and l4

L7          11 L1 AND L2 AND L3 AND L4

=> s l1 and l2 and l3 and l5

L8           4 L1 AND L2 AND L3 AND L5

=> file stnguide

COST IN U.S. DOLLARS                            SINCE FILE      TOTAL
                                                   ENTRY    SESSION
FULL ESTIMATED COST                                 2.85       3.73

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=> file hcaplus
COST IN U.S. DOLLARS                            SINCE FILE      TOTAL
                                                   ENTRY    SESSION
FULL ESTIMATED COST                                 0.07       3.80

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FILE COVERS 1907 - 6 Aug 2009  VOL 151 ISS 6
FILE LAST UPDATED: 4 Aug 2009  (20090804/ED)
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REVISED CLASS FIELDS (/NCL) LAST RELOADED:  Jun 2009
USPTO MANUAL OF CLASSIFICATIONS THESAURUS ISSUE DATE:  Jun 2009

HCAplus now includes complete International Patent Classification (IPC)
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of databases have been updated to include new citing references
information.  This enhancement may impact record import into
database management software.  For additional information, refer
to NEWS 22.

=> s isolation or purification or extraction or separation or isolated or purified

        291282 ISOLATION
        375655 PURIFICATION
        193140 EXTRACTION
        242472 SEPARATION
        939122 ISOLATED
        466181 PURIFIED
L9     2168835 ISOLATION OR PURIFICATION OR EXTRACTION OR SEPARATION OR ISOLATE
               D OR PURIFIED

=> s l1 and l9 and l3

L10         473 L1 AND L9 AND L3

=> s l1 and l9 and l3 and l4

L11          17 L1 AND L9 AND L3 AND L4

=> s l1 and l9 and l3 and l5

L12           5 L1 AND L9 AND L3 AND L5

=> file stnguide

COST IN U.S. DOLLARS                                SINCE FILE      TOTAL
                                                       ENTRY    SESSION
FULL ESTIMATED COST                                     2.85       6.65

FILE 'STNGUIDE' ENTERED AT 16:12:35 ON 06 AUG 2009
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FILE CONTAINS CURRENT INFORMATION.
LAST RELOADED: Jul 31, 2009 (20090731/UP).

=> file hcaplus
COST IN U.S. DOLLARS                                SINCE FILE      TOTAL
                                                       ENTRY    SESSION
FULL ESTIMATED COST                                     0.07       6.72

FILE 'HCAPLUS' ENTERED AT 16:12:49 ON 06 AUG 2009
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FILE COVERS 1907 - 6 Aug 2009 VOL 151 ISS 6
FILE LAST UPDATED: 4 Aug 2009 (20090804/ED)
REVISED CLASS FIELDS (/NCL) LAST RELOADED: Jun 2009
USPTO MANUAL OF CLASSIFICATIONS THESAURUS ISSUE DATE: Jun 2009

HCAplus now includes complete International Patent Classification (IPC)
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This file contains CAS Registry Numbers for easy and accurate
substance identification.

The ALL, BIB, MAX, and STD display formats in the CA/CAplus family
of databases have been updated to include new citing references
information. This enhancement may impact record import into
database management software. For additional information, refer
to NEWS 22.

=> s l11 or l12

L13 19 L11 OR L12

=> file stnguide

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	2.85	9.57

FILE 'STNGUIDE' ENTERED AT 16:12:50 ON 06 AUG 2009
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FILE CONTAINS CURRENT INFORMATION.
LAST RELOADED: Jul 31, 2009 (20090731/UP).

=> d l13 1-19 ti abs bib
YOU HAVE REQUESTED DATA FROM FILE 'HCAPLUS' - CONTINUE? (Y)/N:y

L13 ANSWER 1 OF 19 HCAPLUS COPYRIGHT 2009 ACS on STN
TI Surface mediated self-assembly of solid-phase

nanoparticles for isolation of biomols.
AB Materials and methods for surface mediated self assembly of nanoparticles for the isolation of biomols. is provided.
AN 2009:859079 HCAPLUS <<LOGINID::20090806>>
TI Surface mediated self-assembly of solid-phase nanoparticles for isolation of biomols.
IN Utermohlen, Joseph G.; Hogan, Michael E.; Diggins, Paul E.
PA Argylla Technologies, LLC, USA
SO U.S. Pat. Appl. Publ., 22pp., Cont.-in-part of U.S. Ser. No. 338,124.
CODEN: USXXCO
DT Patent
LA English
FAN.CNT 3

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 20090182120	A1	20090716	US 2008-54325	20080324
	US 20060177855	A1	20060810	US 2006-338124	20060123
PRAI	US 2005-646155P	P	20050121		
	US 2005-701630P	P	20050722		
	US 2006-338124	A2	20060123		
	US 2007-896479P	P	20070322		

L13 ANSWER 2 OF 19 HCAPLUS COPYRIGHT 2009 ACS on STN
TI Method for isolating viral nucleic acids
AB The invention relates to a method for parallel isolation of double- and single-stranded viral nucleic acids from biol. samples without separating of double-and single-stranded nucleic acids. The samples are treated with conventional lysis buffers (high salt concns., or low salt concns. or with proteolytic enzymes). The sample containing nucleic acids before lysis or after lysis or homogenization is adjusted with an acidic binding buffer containing at least one non-ionic detergent in high concentration such that the total nucleic acids are adsorbed on a solid support.
AN 2008:71854 HCAPLUS <<LOGINID::20090806>>
DN 148:114245
TI Method for isolating viral nucleic acids
IN Hillebrand, Timo
PA Aj Innuscreen GmbH, Germany
SO PCT Int. Appl., 18pp.
CODEN: PIXXD2
DT Patent
LA German
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2008006865	A1	20080117	WO 2007-EP57131	20070711
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW			
	RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
	DE 102006032610	A1	20080124	DE 2006-102006032610	20060711
	EP 2041310	A1	20090401	EP 2007-787402	20070711

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        R:  AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE,
            IS, IT, LI, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR,
            AL, BA, HR, MK, RS
PRAI DE 2006-102006032610 A     20060711
     WO 2007-EP57131      W     20070711
RE.CNT  9     THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS RECORD
              ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13  ANSWER 3 OF 19  HCAPLUS  COPYRIGHT 2009 ACS on STN
TI   Solid phase extraction of RNA with
     the use of an alkaline reagent and functionalized magnetic particles
AB   Methods and materials are disclosed for rapid and simple extraction and
     isolation of nucleic acids, particularly RNA, from a
     biol. sample involving the use of an alkaline reagent followed by an acidic
     solution and a solid phase binding material (e.g.,
     magnetic particles functionalized with a tributylphosphonium NAB group and
     a cleavable arylthioester linkage) that has the ability to liberate
     nucleic acids from biol. samples, including whole blood, without first
     performing any preliminary lysis to disrupt cells or viruses.  No
     detergents or chaotropic substances for lysing cells or viruses are needed
     or used.  Viral, bacterial and mammalian genomic RNA can be
     obtained using the method of the invention.  RNA obtained by the
     present method is suitable for use in downstream processes such as RT-PCR.
AN   2007:907280  HCAPLUS <<LOGINID::20090806>>
DN   147:251698
TI   Solid phase extraction of RNA with
     the use of an alkaline reagent and functionalized magnetic particles
IN   Akhavan-Tafti, Hashem; De Silva, Renuka; Eickholt, Robert A.; Mazelis,
     Michael E.; Xie, Wenhuas; Handley, Richard S.; Bray, Monica A.;
     Mastronardi, Michelle L.; O'Conner, Elizabeth A.; Siripurapu, Sarada
PA   Nexgen Diagnostics LLC, USA
SO   U.S. Pat. Appl. Publ., 15pp.
     CODEN: USXXCO
DT   Patent
LA   English
FAN.CNT 1
     PATENT NO.          KIND   DATE        APPLICATION NO.          DATE
     --------------      ----   --------    -------------------      --------
PI   US 20070190526      A1     20070816    US 2007-706547           20070215
     AU 2007217092       A1     20070830    AU 2007-217092           20070216
     CA 2642883          A1     20070830    CA 2007-2642883          20070216
     WO 2007098379       A2     20070830    WO 2007-US62270          20070216
     WO 2007098379       A3     20081120
         W:  AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH,
             CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD,
             GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN,
             KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK,
             MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO,
             RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT,
             TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW
         RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE,
             IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ,
             CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH,
             GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
             KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA
     EP 1989332          A2     20081112    EP 2007-757082           20070216
         R:  AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE,
             IS, IT, LI, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, AL,
             BA, HR, MK, RS
     JP 2009527228       T      20090730    JP 2008-555501           20070216
     IN 2008KN03389      A      20090213    IN 2008-KN3389           20080819
```

	KR 2009003219	A	20090109	KR 2008-722362	20080912
PRAI	US 2006-773881P	P	20060216		
	WO 2007-US62270	W	20070216		

L13 ANSWER 4 OF 19 HCAPLUS COPYRIGHT 2009 ACS on STN

TI Purification of RNA by lysis at acidic pH and capture with immobilized quaternary salts

AB Methods and materials are disclosed for rapid and simple extraction and isolation of RNA from a biol. sample involving the use of an acidic solution and a solid phase binding material that has the ability to liberate nucleic acids from biol. samples, including whole blood, without first performing any preliminary lysis to disrupt cells or viruses. No detergents or chaotropic substances for lysing cells or viruses are needed or used. Materials are lysed at an acidic pH and the liberated RNA is captured by a quaternized salt immobilized on a carrier, such as a magnetic particle. Viral, bacterial and mammalian genomic RNA can be isolated using the method of the invention. RNA isolated by the present method is suitable for use in downstream processes such as RT-PCR. Preparation of the capture moiety 4-(4-HSC6H4SCO)C6H4CH2P+Bu3.I- and its use in recovery of RNA from Escherichia coli, blood, and com. synthetic RNA prepns. is described. The capture moiety may contain a labile group that can be used to release it from the carrier.

AN 2007:874274 HCAPLUS <<LOGINID::20090806>>

DN 147:228308

TI Purification of RNA by lysis at acidic pH and capture with immobilized quaternary salts

IN Akhavan-Tafti, Hashem

PA Nexgen Diagnostics LLC, USA

SO U.S. Pat. Appl. Publ., 15pp., Cont.-in-part of U.S. Provisional Ser. No. 771,510.
CODEN: USXXCO

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 20070185322	A1	20070809	US 2007-703459	20070207
	AU 2007213693	A1	20070816	AU 2007-213693	20070208
	CA 2641615	A1	20070816	CA 2007-2641615	20070208
	WO 2007092916	A3	20071122	WO 2007-US61826	20070208
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW			
	RW: AP, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, EA, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, EP, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, OA, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
	EP 1984383	A2	20081029	EP 2007-763168	20070208
	R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR				
	JP 2009525761	T	20090716	JP 2008-554496	20070208
	IN 2008KN03390	A	20090213	IN 2008-KN3390	20080819
	KR 2009003205	A	20090109	KR 2008-721901	20080908
	CN 101400689	A	20090401	CN 2007-80008550	20080910
PRAI	US 2006-771510P	P	20060208		

US 2007-703459 A 20070207
WO 2007-US61826 W 20070208

L13 ANSWER 5 OF 19 HCAPLUS COPYRIGHT 2009 ACS on STN
TI Methods for nucleic acid extraction and purification
AB The present invention provides methods for extraction and purification of genomic DNA and RNA. Cells are chemical lysed in the presence of a solid phase optionally coated with a charge switch material and nuclear material is flocculated/precipitated Charge switch materials generally comprise a postive charge to bind neg. charged nucleic acids. Genomic DNA can be collected from the precipitate and purified. RNA present in the supernatant can be collected by binding to a solid phase and purified.
AN 2006:31704 HCAPLUS <<LOGINID::20090806>>
DN 144:103508
TI Methods for nucleic acid extraction and purification
IN Baker, Matthew J.; Stevenson, Anthony; Buckels, John
PA Invitrogen Corporation, USA
SO PCT Int. Appl., 43 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2006004611	A2	20060112	WO 2005-US22624	20050627
	WO 2006004611	A3	20060928		
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	US 20060024712	A1	20060202	US 2005-167907	20050627
PRAI	GB 2004-14302	A	20040625		
	US 2004-582879P	P	20040625		
	GB 2004-22299	A	20041007		

L13 ANSWER 6 OF 19 HCAPLUS COPYRIGHT 2009 ACS on STN
TI Purification of nucleic acids from solutions by reversible binding of polynucleotides to the surface of a magnetic microparticle
AB The invention relates to methods of separating polynucleotides, such as DNA, RNA and PNA, from solns. containing polynucleotides by reversibly binding the polynucleotides to a solid surface, such as a magnetic microparticle. The invention allows to obtain polynucleotides sufficiently free from contaminants for mol. biol. applications from animal tissues and body fluids.
AN 2005:1028001 HCAPLUS <<LOGINID::20090806>>
DN 143:300280
TI Purification of nucleic acids from solutions by reversible binding of polynucleotides to the surface of a magnetic microparticle
IN Latham, Gary J.; Fang, Xingwang; Conrad, Richard C.; Kemppainen, Jon A.; Setterquist, Robert A.; Pasloske, Brittan L.
PA Ambion, Inc., USA

SO U.S. Pat. Appl. Publ., 36 pp.
CODEN: USXXCO
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 20050208510	A1	20050922	US 2004-955974	20040930
	WO 2005089929	A2	20050929	WO 2005-US9189	20050318
	WO 2005089929	A3	20051124		
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	RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, MR, NE, SN, TD, TG				
	EP 1735466	A2	20061227	EP 2005-760857	20050318
	R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR				
	JP 2007529229	T	20071025	JP 2007-504164	20050318
PRAI	US 2004-554278P	P	20040318		
	US 2004-955974	A2	20040930		
	WO 2005-US9189	W	20050318		

OSC.G 1 THERE ARE 1 CAPLUS RECORDS THAT CITE THIS RECORD (1 CITINGS)

L13 ANSWER 7 OF 19 HCAPLUS COPYRIGHT 2009 ACS on STN
TI Methods for separating unincorporated deoxyribonucleotide triphosphates or salts from DNA or purification of other analytes using coated magnetic hydroxylapatite beads
AB The present invention provides a material for separating an analyte from an undesired constituent, which material comprises a solid phase and a coating, wherein the solid phase is capable of binding the undesired constituent, and wherein the coating covers the exposed surface of the solid phase to an extent that any binding of the solid phase to the analyte is impeded. In particular, it provides methods for separating unincorporated deoxyribonucleotide triphosphates from DNA or purification of other analytes prior to anal. using coated magnetic hydroxylapatite beads.
AN 2005:121095 HCAPLUS <<LOGINID::20090806>>
DN 142:172863
TI Methods for separating unincorporated deoxyribonucleotide triphosphates or salts from DNA or purification of other analytes using coated magnetic hydroxylapatite beads
IN Goldsborough, Andrew
PA Cyclops Genome Sciences Limited, UK
SO PCT Int. Appl., 42 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2005012522	A1	20050210	WO 2004-GB3201	20040723
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,				

LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI,
NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY,
TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM,
AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK,
EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE,
SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE,
SN, TD, TG

EP	1649016	A1	20060426	EP 2004-743533	20040723
R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, FI, RO, CY, TR, BG, CZ, EE, HU, PL, SK				
JP	2006527993	T	20061214	JP 2006-520900	20040723
US	20080220413	A1	20080911	US 2008-565694	20080324
PRAI GB	2003-17199	A	20030723		
GB	2003-19422	A	20030819		
WO	2004-GB3201	W	20040723		

L13 ANSWER 8 OF 19 HCAPLUS COPYRIGHT 2009 ACS on STN
TI Locked nucleic acid capture probes for isolation of homopolymeric nucleotide sequences and use in diagnosis of viral infections in humans
AB This invention presents methods and use of locked nucleic acid capture probes for isolation of homopolymeric nucleotide sequences and use in diagnosis of viral infections in humans. A method for isolating nucleic acid mols. having a repeating nucleotide sequence or a homopolymeric nucleotide sequence, e.g. a poly A stretch, is described. In particular, the method uses oligomeric capture probes spiked with various amts. of locked nucleic acid (LNA). The invention further describes methods for the isolation of RNA mols., for example polyadenylated mRNA mols., which overcome the problems of rapid RNA degradation during isolation and anal. of such nucleic acid mols. This is of major clin. and diagnostic importance, especially when dealing with RNA viruses, such as retroviruses or when analyzing rare or low-abundant mRNAs or mRNAs from biopsies or tissues enriched with RNases.
AN 2004:203927 HCAPLUS <<LOGINID::20090806>>
DN 140:265567
TI Locked nucleic acid capture probes for isolation of homopolymeric nucleotide sequences and use in diagnosis of viral infections in humans
IN Kauppinen, Sakari; Jacobsen, Nana
PA Exiqon A/S, Den.
SO PCT Int. Appl., 104 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2004020575	A2	20040311	WO 2003-IB6354	20030620
WO 2004020575	A3	20041223		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				

	AU 2003288474	A1	20040319	AU 2003-288474	20030620
	US 20050053942	A1	20050310	US 2003-601140	20030620
	EP 1527175	A2	20050504	EP 2003-780549	20030620
	EP 1527175	B1	20090527		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
	AT 432347	T	20090615	AT 2003-780549	20030620
	US 20080096191	A1	20080424	US 2007-893108	20070814
PRAI	US 2002-390928P	P	20020624		
	US 2003-601140	B1	20030620		
	WO 2003-IB6354	W	20030620		

OSC.G 1 THERE ARE 1 CAPLUS RECORDS THAT CITE THIS RECORD (1 CITINGS)
RE.CNT 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13 ANSWER 9 OF 19 HCAPLUS COPYRIGHT 2009 ACS on STN
TI Compositions and methods for using a solid support to purify RNA
AB The invention concerns a method for purifying substantially pure and undegraded RNA from biol. material comprising RNA, comprising the steps of: (a) mixing the biol. material with an RNA Lysing/ Binding Solution buffered at a pH of greater than about 7, the RNA Lysing/Binding Solution comprising an RNA-complexing salt; (b) contacting the mixture to a solid support such that nucleic acids comprising substantially undegraded RNA in the mixture preferentially bind to the solid support; (c) washing the solid support with a series of RNA wash solns. to remove biol. materials other than bound nucleic acids comprising substantially undegraded RNA, wherein the series of wash solns. comprises a first wash comprising alc. and an RNA-complexing salt at a concentration of at least 1 M and a second wash comprising an alc., buffer and an optional chelator; and (d) preferentially eluting the bound substantially undegraded RNA from the solid support with an RNA Elution Solution in order to obtain substantially pure and undegraded RNA. Reagents, methods and kits for the purification of RNA from biol. materials are provided.
AN 2004:80382 HCAPLUS <<LOGINID::20090806>>
DN 140:107795
TI Compositions and methods for using a solid support to purify RNA
IN Bair, Robert Jackson; Heath, Ellen M.; Meehan, Heather; Paulsen, Kim Elayne; Wages, John M.
PA USA
SO U.S. Pat. Appl. Publ., 19 pp., Cont.-in-part of U.S. Ser. No. 974,798.
CODEN: USXXCO
DT Patent
LA English
FAN.CNT 3

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 20040019196	A1	20040129	US 2003-418194	20030416
	US 7148343	B2	20061212		
	US 20030073830	A1	20030417	US 2001-974798	20011012
	CA 2463317	A1	20030424	CA 2001-2463317	20011012
	AU 2002211719	A1	20030428	AU 2002-211719	20011012
	AU 2002211719	B2	20070614		
	EP 1438426	A1	20040721	EP 2001-979794	20011012
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
	JP 2005505305	T	20050224	JP 2003-536461	20011012

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     JP 3979996           B2    20070919
     AU 2004233035        A1    20041104    AU 2004-233035         20040415
     CA 2522446           A1    20041104    CA 2004-2522446        20040415
     WO 2004094635        A2    20041104    WO 2004-US12033        20040415
     WO 2004094635        A3    20041216
          W:  AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH,
              CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD,
              GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,
              LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI,
              NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY,
              TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
          RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ,
              BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE,
              ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI,
              SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN,
              TD, TG
     EP 1618194           A2    20060125    EP 2004-760008         20040415
          R:  AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
              IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, PL, SK, HR
     JP 2006523463        T     20061019    JP 2006-513124         20040415
     US 20050032105       A1    20050210    US 2004-909724         20040802
     US 20070043216       A1    20070222    US 2006-589364         20061030
PRAI US 2001-974798       A2    20011012
     WO 2001-US32073      W     20011012
     US 2003-418194       A     20030416
     WO 2004-US12033      W     20040415
OSC.G   1     THERE ARE 1 CAPLUS RECORDS THAT CITE THIS RECORD (1 CITINGS)
RE.CNT  56    THERE ARE 56 CITED REFERENCES AVAILABLE FOR THIS RECORD
              ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13  ANSWER 10 OF 19  HCAPLUS  COPYRIGHT 2009 ACS on STN
TI   Methods and kits for isolating nucleic acids from biological samples using
     hydrophilic magnetic particles and chaotrope agents
AB   A process for isolating nucleic acids using hydrophilic magnetic particles
     is provided.  The nucleic acid in biol. sample is bound to the magnetic
     particles in the presence of the chaotrope.  The methods also comprises
     contacting the sample with the nucleic acid binding solid
     phase in the presence of a liquid phase comprising the chaotrope.
     The methods also comprises optionally separating the solid
     phase with the nucleic acid bound thereto from the liquid phase,
     wherein the solid phase bears acid groups on its
     surface.  The chaotrope comprises a guanidinium salt, urea, or an iodide,
     chlorate, perchlorate or (iso)thiocyanate.
AN   2003:913301  HCAPLUS <<LOGINID::20090806>>
DN   139:377555
TI   Methods and kits for isolating nucleic acids from biological samples using
     hydrophilic magnetic particles and chaotrope agents
IN   Deggerdal, Arne; Skagestad, Vidar
PA   Qiagen A/S, Norway
SO   PCT Int. Appl., 19 pp.
     CODEN: PIXXD2
DT   Patent
LA   English
FAN.CNT 1
     PATENT NO.          KIND   DATE        APPLICATION NO.        DATE
     --------------      ----   --------    -------------------    --------
PI   WO 2003095646        A1    20031120    WO 2003-IB1822         20030509
          W:  AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
              CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
              GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
              LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH,
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PL, PT, RO, RU, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

AU 2003230070 A1 20031111 AU 2003-230070 20030509
PRAI GB 2002-10766 A 20020510
WO 2003-IB1822 W 20030509
RE.CNT 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13 ANSWER 11 OF 19 HCAPLUS COPYRIGHT 2009 ACS on STN
TI Methods, reagents and kits for isolating RNA from environmental or biological samples
AB Reagents, methods and kits for the purification of RNA from biol. or environmental samples are provided. The method comprises mixing said material with an RNA binding solution buffered at a pH of greater than 7 wherein the RNA binding solution comprises an RNA complexing salt from from strong chaotropic agents. RNA is bound to non-silica solid support selected from cellulose, cellulose acetate, nitrocellulose, nylon, polyester, polyethersulfone, polyolefin, or polyvinylidene fluoride. The non-silica solid support is contained in a vessel such as centrifuge tubes, spin tubes, syringes, cartridges, chambers, multiple well plates and test tubes.
AN 2003:300642 HCAPLUS <<LOGINID::20090806>>
DN 138:317132
TI Methods, reagents and kits for isolating RNA from environmental or biological samples
IN Heath, Ellen M.; Wages, John M.
PA USA
SO U.S. Pat. Appl. Publ., 14 pp.
CODEN: USXXCO
DT Patent
LA English
FAN.CNT 3

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI US 20030073830	A1	20030417	US 2001-974798	20011012
CA 2463317	A1	20030424	CA 2001-2463317	20011012
WO 2003033739	A1	20030424	WO 2001-US32073	20011012
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
AU 2002211719	A1	20030428	AU 2002-211719	20011012
AU 2002211719	B2	20070614		
EP 1438426	A1	20040721	EP 2001-979794	20011012
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
JP 2005505305	T	20050224	JP 2003-536461	20011012
JP 3979996	B2	20070919		
US 20040019196	A1	20040129	US 2003-418194	20030416
US 7148343	B2	20061212		

	US 20050032105	A1	20050210	US 2004-909724	20040802
	US 20070043216	A1	20070222	US 2006-589364	20061030
PRAI	US 2001-974798	A	20011012		
	WO 2001-US32073	W	20011012		
	US 2003-418194	A2	20030416		

OSC.G 1 THERE ARE 1 CAPLUS RECORDS THAT CITE THIS RECORD (1 CITINGS)

L13 ANSWER 12 OF 19 HCAPLUS COPYRIGHT 2009 ACS on STN

TI Methods and kits for the purification of nucleic acids from bacterial cells using a single reagent containing polyethylene glycol and binding to paramagnetic beads

AB The invention includes reagents and methods for the isolation of nucleic acids. The reagents described herein contain a nucleic acid precipitating agent and a solid phase carrier. The reagents can optionally be formulated to cause the lysis of a cell. These reagents can be used to isolate a target nucleic acid mol. from a cell or a solution containing a mixture of different size nucleic acid mols. In a preferred embodiment plasmid DNA from bacterial cells are purified by precipitation with 1-4% polyethylene glycol (mol. weight of 8000) and 0.5M salt concentration The DNA is further purified by reversible binding to paramagnetic beads that are coated with amine or encapsulated carboxyl groups. The first reagent allows purification of DNA greater than 10 kb, while a second round of purification allows purification of DNA greater than 2.4 kb from a mixture of nucleic acids 7% polyethylene glycol. Magnetic fields of about 1000 G are applied to the wells of a microtiter plate using a magnetic plate holder containing an N35 magnet for removal of paramagnetic beads following DNA purification The disclosed reagents and methods provides a simple, robust and readily automatable means of nucleic acid isolation and purification which produces high quality nucleic acid mols. suitable for: capillary electrophoresis, nucleotide sequencing, reverse transcription cloning the transfection, transduction or microinjection of mammalian cells, gene therapy protocols, the in vitro synthesis of RNA probes, cDNA library construction and PCR amplification.

AN 2002:539860 HCAPLUS <<LOGINID::20090806>>

DN 137:89428

TI Methods and kits for the purification of nucleic acids from bacterial cells using a single reagent containing polyethylene glycol and binding to paramagnetic beads

IN McKernan, Kevin J.

PA Whitehead Institute for Biomedical Research, USA

SO PCT Int. Appl., 45 pp.
CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2002055727	A2	20020718	WO 2002-US353	20020109
	WO 2002055727	A3	20021003		
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				

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     CA 2433746          A1    20020718    CA 2002-2433746        20020109
     AU 2002239826       A1    20020724    AU 2002-239826         20020109
     US 20020106686      A1    20020808    US 2002-42923          20020109
     EP 1349951          A2    20031008    EP 2002-705692         20020109
        R:  AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
            IE, SI, LT, LV, FI, RO, MK, CY, AL, TR
     US 20060024701      A1    20060202    US 2005-126775         20050511
PRAI US 2001-260774P     P     20010109
     US 2002-42923       B1    20020109
     WO 2002-US353       W     20020109
OSC.G   5     THERE ARE 5 CAPLUS RECORDS THAT CITE THIS RECORD (5 CITINGS)
RE.CNT  4     THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD
              ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13  ANSWER 13 OF 19  HCAPLUS  COPYRIGHT 2009 ACS on STN
TI   Methods and kits for isolating nucleic acids from leukocytes by binding to
     antibodies on a solid support
AB   The present invention relates to a method of isolating nucleic acid from a
     blood sample.  The method involves selectively isolating leukocytes from
     said sample by binding said leukocytes to a solid
     support containing a binding partner specific for the leukocyte, for
     example an antibody.  The antibody can bind an antigen selected from one
     of more of the following: HLA-I, CD11a, CD18, CD45, CD46, CD50, CD82,
     CD162, CD5 and CD15 and a specific example shows a combination of CD45 and
     CD15.  The said leukocytes are lysed in detergents to release nucleic
     acids which are subsequently bound to a second solid
     support which is neg. charged.  Kits for isolating nucleic acid
     from samples form further embodiments of the invention.
AN   2001:904506  HCAPLUS <<LOGINID::20090806>>
DN   136:15912
TI   Methods and kits for isolating nucleic acids from leukocytes by binding to
     antibodies on a solid support
IN   Bergholtz, Stine; Korsnes, Lars; Andreassen, Jack
PA   Dynal Biotech Asa, Norway; Jones, Elizabeth Louise
SO   PCT Int. Appl., 51 pp.
     CODEN: PIXXD2
DT   Patent
LA   English
FAN.CNT 1
     PATENT NO.          KIND   DATE        APPLICATION NO.        DATE
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PI   WO 2001094572       A1     20011213    WO 2001-GB2472         20010605
        W:  AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
            CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
            GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
            LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT,
            RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US,
            UZ, VN, YU, ZA, ZW
        RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
            DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF,
            BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
     CA 2410888          A1     20011213    CA 2001-2410888        20010605
     CA 2410888          C      20080916
     EP 1290155          A1     20030312    EP 2001-934205         20010605
     EP 1290155          B1     20060809
        R:  AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
            IE, SI, LT, LV, FI, RO, MK, CY, AL, TR
     AU 2001260507       B2     20060831    AU 2001-260507         20010605
     AT 335815           T      20060915    AT 2001-934205         20010605
     ES 2269399          T3     20070401    ES 2001-934205         20010605
     US 20030180754      A1     20030925    US 2003-297301         20030430
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     US 20080293035       A1    20081127    US 2008-98411          20080404
PRAI GB 2000-13658        A     20000605
     WO 2001-GB2472       W     20010605
     US 2003-297301       B1    20030430
OSC.G   2     THERE ARE 2 CAPLUS RECORDS THAT CITE THIS RECORD (2 CITINGS)
RE.CNT  4     THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD
              ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13  ANSWER 14 OF 19  HCAPLUS  COPYRIGHT 2009 ACS on STN
TI   Methods for detecting and measuring spliced nucleic acids and method of
     cytoplasmic nucleic acid preparation
AB   A simplified method for preparing a biol. sample to release cytoplasmic
     nucleic acid, preferably spliced mRNA, suitable for amplification, while
     minimizing the release of nuclear genetic material is disclosed.  A buffer
     containing a soluble salt with ionic strength of particular range and a
non-ionic
     detergent are used to lyse the cells.  MRNA is then purified by
     contacting the sample with a solid support joined to
     an immobilized oligonucleotide which would form stable hybridization
     complex with the mRNA.  Immobilized oligonucleotide preferably contains a
     poly-T sequence.  A method of detecting and measuring the amount of fusion
     nucleic acid, notably spliced mRNA present in the sample, following
     nucleic acid amplification, is also disclosed.  A fusion nucleic acid to
     be detected contain a splice junction site, and primers designed to have
     sequences complementary to and around the splice-junction site are used to
     amplify the nucleic acid.  The amplified nucleic acid strand is detected
     with an oligonucleotide probe which specifically hybridizes to the
     amplified strand.  Nucleic acid of chronic myelogenous leukemia patient
     and that resulting from bcr-abl translocation were detected by the method.
AN   2000:85055  HCAPLUS <<LOGINID::20090806>>
DN   132:147583
TI   Methods for detecting and measuring spliced nucleic acids and method of
     cytoplasmic nucleic acid preparation
IN   Harvey, Richard C.; Eastman, Paul S.
PA   Gen-Probe Incorporated, USA
SO   PCT Int. Appl., 52 pp.
     CODEN: PIXXD2
DT   Patent
LA   English
FAN.CNT 1
     PATENT NO.          KIND   DATE        APPLICATION NO.        DATE
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PI   WO 2000005418       A1     20000203    WO 1999-US16832        19990723
         W:  AU, CA, JP
         RW: AT, BE, CH, DE, DK, ES, FR, GB, IT, LU, NL, SE
     US 6849400           B1     20050201    US 1998-121239         19980723
     CA 2337106           A1     20000203    CA 1999-2337106        19990723
     AU 9951288           A      20000214    AU 1999-51288          19990723
     AU 767568            B2     20031113
     EP 1109932           A1     20010627    EP 1999-935912         19990723
     EP 1109932           B1     20040616
         R:  AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, FI
     JP 2002521037        T      20020716    JP 2000-561364         19990723
     AT 269417            T      20040715    AT 1999-935912         19990723
     ES 2221750           T3     20050101    ES 1999-935912         19990723
PRAI US 1998-121239       A      19980723
     US 1997-53509P       P      19970723
     WO 1999-US16832      W      19990723
OSC.G   3     THERE ARE 3 CAPLUS RECORDS THAT CITE THIS RECORD (3 CITINGS)
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L13 ANSWER 15 OF 19 HCAPLUS COPYRIGHT 2009 ACS on STN
TI Solid phase technique for selectively isolating nucleic acids
AB A method of isolating target nucleic acid mols. from a solution comprising a mixture of different size nucleic acid mols., in the presence or absence of other biomols., by selectively facilitating the adsorption of a particular species of nucleic acid mol. to the functional group-coated surface of magnetically responsive paramagnetic microparticles is disclosed. Separation is accomplished by manipulating the ionic strength and polyalkylene glycol concentration of the solution to selectively precipitate, and reversibly adsorb, the target species of nucleic acid mol., characterized by a particular mol. size, to paramagnetic microparticles, the surfaces of which act as a bioaffinity adsorbent for the nucleic acids. The target nucleic acid is isolated from the starting mixture based on mol. size and through the removal of magnetic beads to which the target nucleic acid mols. have been adsorbed. The disclosed method provides a simple, robust and readily automatable means of nucleic acid isolation and purification which produces high quality nucleic acid mols. suitable for: capillary electrophoresis, nucleotide sequencing, reverse transcription cloning the transfection, transduction or microinjection of mammalian cells, gene therapy protocols, the in vitro synthesis of RNA probes, cDNA library construction and PCR amplification.
AN 1999:736906 HCAPLUS <<LOGINID::20090806>>
DN 131:334336
TI Solid phase technique for selectively isolating nucleic acids
IN McKernan, Kevin; McEwan, Paul; Morrison, William
PA Whitehead Institute for Biomedical Research, USA
SO PCT Int. Appl., 46 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9958664	A1	19991118	WO 1999-US10572	19990513
	W: CA, JP				
	RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
	US 6534262	B1	20030318	US 1999-311317	19990513
	US 20030235839	A1	20031225	US 2003-346714	20030116
	US 20040214175	A9	20041028		
	US 20060003357	A1	20060105	US 2005-129218	20050513
PRAI	US 1998-85480P	P	19980514		
	US 1999-121779P	P	19990226		
	US 1999-311317	A1	19990513		
	US 2003-346714	A3	20030116		

OSC.G 11 THERE ARE 11 CAPLUS RECORDS THAT CITE THIS RECORD (12 CITINGS)

L13 ANSWER 16 OF 19 HCAPLUS COPYRIGHT 2009 ACS on STN
TI Methods and compositions for isolating nucleic acids
AB Compns. and methods are disclosed for isolating nucleic acids from biol. tissues and cells (including tumor cells) and for tissue/cell solubilization for other mol. biol. uses, wherein the compns. comprise, in part, novel combinations of chaotropic agents and aromatic alcs. which act synergistically to effect better tissue/protein solubilization. The inventive compns. further include aprotic solvents for deactivation of RNases and denaturization of proteins, as well as detergents for enhancing cell lysis and nucleoprotein dissociation The inventive methods also comprise the use of a centrifuge, a solid-support matrix, and a

microporous membrane for final isolation of the precipitated nucleic acids, resulting in high yield and purity of the precipitated nucleic acid.

AN 1997:400479 HCAPLUS <<LOGINID::20090806>>
DN 127:78238
OREF 127:14901a,14904a
TI Methods and compositions for isolating nucleic acids
IN Wiggins, James C.
PA USA
SO U.S., 15 pp.
CODEN: USXXAM
DT Patent
LA English
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI US 5637687	A	19970610	US 1993-115184	19930831
PRAI US 1993-115184		19930831		

OSC.G 10 THERE ARE 10 CAPLUS RECORDS THAT CITE THIS RECORD (10 CITINGS)
RE.CNT 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13 ANSWER 17 OF 19 HCAPLUS COPYRIGHT 2009 ACS on STN
TI Isolation of nucleic acid from biological sample, method comprising nucleic acid binding to solid support then separation from support, and kit comprising detergents and other components
AB The present invention provides a method of isolating nucleic acid from a sample, said method comprising contacting said sample with a detergent and a solid support, whereby soluble nucleic acid in said sample is bound to the support, and separating said support with bound nucleic acid from the sample. Where the method of the invention is used to isolate DNA, it may conveniently be coupled with a further step to isolate RNA from the same sample.
AN 1996:458048 HCAPLUS <<LOGINID::20090806>>
DN 125:107039
OREF 125:19863a,19866a
TI Isolation of nucleic acid from biological sample, method comprising nucleic acid binding to solid support then separation from support, and kit comprising detergents and other components
IN Deggerdal, Arne Helge; Larsen, Frank
PA Dynal A/s, Norway; Dzieglewska, Hanna Eva
SO PCT Int. Appl., 53 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 9618731	A2	19960620	WO 1995-GB2893	19951212
WO 9618731	A3	19960912		
W: AL, AM, AT, AU, BB, BG, BR, BY, CA, CH, CN, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IS, JP, KE, KG, KP, KR, KZ, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK				
RW: KE, LS, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG				
CA 2207608	A1	19960620	CA 1995-2207608	19951212
CA 2207608	C	20090407		
AU 9641829	A	19960703	AU 1996-41829	19951212

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     AU 706211            B2    19990610
     EP 796327            A2    19970924    EP 1995-940351         19951212
     EP 796327            B1    20040728
        R:  AT, BE, CH, DE, FR, GB, IT, LI, SE
     JP 11501504          T     19990209    JP 1996-518463         19951212
     JP 3787354           B2    20060621
     AT 272110            T     20040815    AT 1995-940351         19951212
     US 20040215011       A1    20041028    US 1997-849686         19970821
     US 20060058519       A1    20060316    US 2005-234001         20050923
     US 7173124           B2    20070206
     US 20070190559       A1    20070816    US 2007-671426         20070205
     US 20080300396       A1    20081204    US 2008-54332          20080324
     US 20090068724       A1    20090312    US 2008-130926         20080530
     US 20090149646       A1    20090611    US 2008-130959         20080530
PRAI GB 1994-25138        A     19941212
     WO 1995-GB2893       W     19951212
     US 1997-849686       A1    19970821
     US 2005-234001       A1    20050923
     US 2007-671426       B1    20070205
     US 2008-54332        A1    20080324
OSC.G   18    THERE ARE 18 CAPLUS RECORDS THAT CITE THIS RECORD (19 CITINGS)
RE.CNT  1     THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD
              ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13  ANSWER 18 OF 19  HCAPLUS  COPYRIGHT 2009 ACS on STN
TI   Purification of nucleic acids from solution without
     precipitation by binding to a solid phase
AB   A method of separating polynucleotides, such as DNA, RNA and PNA,
     from solution by reversibly and non-specifically binding them to a solid
     surface, such as a magnetic microparticle, with a functional group-coated
     surface is disclosed.  The salt and polyalkylene glycol concentration of the
     solution is adjusted to levels which result in polynucleotide binding to the
     magnetic microparticles.  The magnetic microparticles with bound
     polynucleotides are separated from the solution and the polynucleotides are
     eluted from the magnetic microparticles.  The method is generally
     applicable to large and small nucleic acids and works with crude prepns.
     such as cleared lysates.  Material can be selectively eluted from the
     particles by controlling the ionic strength of the elution buffer.
AN   1996:350414  HCAPLUS <<LOGINID::20090806>>
DN   125:5056
OREF 125:1147a,1150a
TI   Purification of nucleic acids from solution without
     precipitation by binding to a solid phase
IN   Hawkins, Trevor
PA   Whitehead Institute for Biomedical Research, USA
SO   PCT Int. Appl., 38 pp.
     CODEN: PIXXD2
DT   Patent
LA   English
FAN.CNT 1
     PATENT NO.          KIND   DATE        APPLICATION NO.        DATE
     --------------      ----   --------    -------------------    --------
PI   WO 9609379          A1     19960328    WO 1995-US11839        19950919
        W:  CA, JP
        RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE
     US 5705628          A      19980106    US 1994-309267         19940920
     IL 115352           A      20090211    IL 1995-115352         19950919
     US 5898071          A      19990427    US 1998-2412           19980102
PRAI US 1994-309267      A      19940920
OSC.G   29    THERE ARE 29 CAPLUS RECORDS THAT CITE THIS RECORD (32 CITINGS)
RE.CNT  1     THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD
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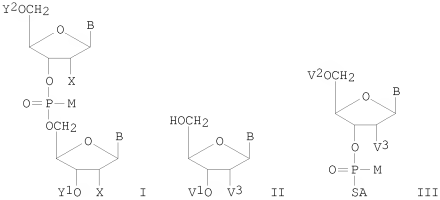
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13 ANSWER 19 OF 19 HCAPLUS COPYRIGHT 2009 ACS on STN
TI Pentavalent synthesis of oligonucleotides containing stereospecific alkylphosphonates and arylphosphonates
GI



AB The present invention provides a method for making R stereospecific alkyl- and aryl-phosphonate linkages between nucleotides. These methods can be used for automated synthesis of oligonucleotides having sequential R stereospecific alkyl- and aryl-phosphonate linkages. The present invention is also directed to the oligonucleotides having several sequential R phosphonate linkages which were produced by the subject methods. Moreover, the present invention provides methods for using the subject oligonucleotides, including methods for regulating the biosynthesis of a DNA, and RNA or a protein and methods for detecting and isolating complementary nucleic acid targets. Title oligonucleotides [I; Y1 = H, phosphate, V1; Y2 = H, phosphate, V2; X = OH, V3; M = alkyl, cycloalkyl, thioxo, etc.; B = (un)substituted purine or pyrimidine residue; V1 = protecting group, solid support , or phosphate attached to the penultimate nucleotide of said oligonucleotide; V2 = protecting group; V3 = H, O-Y3; Y3 = alkyl protecting group; A = activating group] and their intermediates are prepared E.g., 5'-(dimethoxytrityl)thymidyl 3'-methylphosphonoamidate was protected by cyanoethylation in the presence of 4-(N,N-diethylamino)pyridine and (CF3CO)2O at room temperature to give 5'-(dimethoxytrityl)thymidyl 3'-[2-cyanoethyl methylphosphonate], whose oxidation with sulfur (S8) in the presence of MeCN gave the diastereomers of 5'-(dimethoxytrityl)thymidyl 3'-[2-cyanoethyl methylphosphonothioate], which were separated and purified by HPLC; cyanoethyl groups were removed with concentrated NH4OH in EtOH, the deprotected diastereomers were then purified by silica HPLC and the ammonium cation was replaced with Li+ by using a Dowex 50W + 2 exchange column to yield the lithium salts of sep. Sp- and Rp-stereoisomers of 5'-(dimethoxytrityl)thymidyl 3'-methylphosphonothioate. Sp- and Rp-stereoisomers prepared as above were stable and were separated by ion exchange chromatog. or by HPLC using anhydrous or aqueous solvents. The Sp-stereoisomer is reacted with an activator, e.g., 2-chloro-N-methylpyridinium, the intermediate (with retention of configuration) then undergoes an SN2 replacement reaction with a 5'-unprotected nucleoside to give the Rp-configurated dinucleotide; the

displaced 2-thio-N-methylpyridinium mol. is stabilized by resonance tautomerization and does not react with the phosphorus to cause epimerization of the R configuration. A compartmentalized kit for producing a polynucleotide chain of an oligonucleotide having at least 5 sequential R-alkylphosphonate or R-arylphosphonate linkages are claimed. These methods may be used in correcting genetic disorders, e.g., Alzheimer's disease, by inhibiting the production of mutants or over-produced proteins.

AN 1994:409934 HCAPLUS <<LOGINID::20090806>>
DN 121:9934
OREF 121:2104h,2105a
TI Pentavalent synthesis of oligonucleotides containing stereospecific alkylphosphonates and arylphosphonates
IN Wickstrom, Eric; Lebedev, Alexander V.
PA Research Corp. Technologies, Inc., USA
SO PCT Int. Appl., 233 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9400473	A2	19940106	WO 1993-US6277	19930630
	WO 9400473	A3	19940217		
	W: AU, CA, JP, KR				
	RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
	AU 9346611	A	19940124	AU 1993-46611	19930630
PRAI	US 1992-907771	A	19920630		
	WO 1993-US6277	A	19930630		

OS MARPAT 121:9934
RE.CNT 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT